# PCT WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



(51) International Patent Classification 6:	TED (	INDER THE PATENT COOPERATION TREATY (PCT)
	A1	(11) International Publication Number: WO 99/21916
C08K 3/30, 3/34, C08L 23/12, F16L 58/16	AI	(43) International Publication Date: 6 May 1999 (06.05.99
(21) International Application Number: PCT/EP9 (22) International Filing Date: 20 October 1998 (3)		European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR
(22) International Filing Date: 20 October 1998 (2	20.10.9	05, 0K, 15, 11, 50, MC, NE, F1, 35).
(30) Priority Data: MI97A002414 28 October 1997 (28.10.97)		Published T With international search report.
(71) Applicant (for all designated States except US): MG NORTH AMERICA INC. [US/US]; Three Little F tre, 2801 Centerville Road, Wilmington, DE 198 (US).	alls Ce	1-
(72) Inventor; and (75) Inventor/Applicant (for US only): RIGOSI, Gia [IT/IT]; Via Pomposa, 40, I-44100 Ferrara (IT).	ın, Lu	gi
(74) Agent: ZANOLI, Enrico; Montell Italia S.p.A., Pa Trademarks Intellectual Property, Via Pergolesi, 25 Milano (IT).		
(54) Title: POLYPROPYLENE COMPOSITIONS WIT COATING METAL PIPES	НА	HIGH CONTENT OF HEAVY MINERAL FILLERS SUITED FOR
(57) Abstract		
than or equal to 3; (B) 0.1-4 % of a filler compatibilizer;	; and (	0 % of at least one heavy mineral filler having a specific weight greate c) 10-40 % of a heterophasic polyolefin composition containing rubbeing at break greater than 100 to 800 %, and flexural modulus ranging
·		

### FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
СМ	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
cz	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

# "POLYPROPYLENE COMPOSITIONS WITH A HIGH CONTENT OF HEAVY MINERAL FILLERS SUITED FOR COATING METAL PIPES"

The present invention concerns polyolefin compositions with a high content of heavy mineral fillers, particularly suited for coating metal pipes by way of extrusion coating processes.

It is known that polyolefins, and particularly propylene polymers, can be used satisfactorily to coat the outside surface of metal pipes to be used for pipelines designed for liquid and gaseous fluids.

In fact, propylene polymers allow one to obtain coatings with high corrosion resistance and optimum mechanical properties.

However, in cases where the pipes thus coated are to be used for marine pipelines there is often the need for additional external layers of heavy materials, particularly concrete, in order to keep said pipelines from floating.

For example, polyolefin compositions, optionally containing elastomeric polymers, having a high specific weight are described in European patent application EP-A-673966. The polyolefin compositions described in the above mentioned patent application are suitable for coating displaying a good balance of weight, corrosion resistance, and high mechanical properties, rendering unnecessary the further addition of layers of other heavy materials, such as concrete, that are subject to breaking and at times are difficult to apply to the pipes.

The need has now been found for materials that compared to the materials already known provide better mechanical properties while having high weight and being corrosion resistant. In particular, the need is for a material with high elongation at break, good elasticity and impact strength.

WO 99/21916 PCT/EP98/0662<del>5</del>

In response to the above mentioned needs, the Applicant perfected polyolefin compositions having thermoplastic and elastomeric properties, containing high quantities of heavy mineral fillers, easy to apply to the outside surface of metal pipes, allowing the application of single or multiple layers, and possessing the above mentioned balance of properties.

The multi-layer coatings can comprise layers of plastic materials with different properties, such as expanded plastics.

It has been noticed that in order to obtain the required results the compositions must contain a limited maximum quantity of an elastomeric copolymer in order for the compositions to adhere to the pipe to which they are applied; on the other hand, if the quantity of elastomeric copolymer is too low, the desired balance of mechanical properties is not obtained.

Therefore, object of the present invention are polyolefin compositions comprising:

- (A) 60%-90%, preferably 65%-90%, more preferably 75%-85% by weight of at least one heavy mineral filler having a specific weight greater than or equal to 3, such as BaSO<sub>4</sub>, or zirconium silicates;
- (B) 10-40%, preferably 10-35%, more preferably 15-25% by weight of a heterophasic polyolefin composition (having thermoplastic and elastomeric properties); and
- (C) a compatibilizer for fillers in quantities ranging from 0.1% to 4%, preferably 0.2%-2% by weight with respect to the weight of components (A)+(B);

said composition (B) comprising (weight percentage):

30-60% of a propylene homopolymer fraction, said fraction being more that 80% insoluble in xylene at ambient temperature, or a copolymer of propylene with ethylene and/or a C<sub>4</sub>-C<sub>10</sub> α-olefin, containing more than 80% propylene, and being more than 80% insoluble in xylene at ambient temperature (Fraction 1°);

d.

40-70% of a fraction of a copolymer of ethylene with propylene and/or a C<sub>4</sub>-C<sub>10</sub> α-olefin, and optionally minor quantities of a diene; said copolymer containing from 15 to 70% of ethylene, and being soluble in xylene at ambient temperature (Fraction 2°); and

3) 0-30%, with respect to the sum of Fractions (1°) and (2°), of a copolymer fraction containing ethylene, said fraction being insoluble in xylene at ambient temperature (Fraction 3°).

Said compositions have a flow index (ASTM D 1238, condition L, MIL) ranging from 0.5 to 10 g/10 minutes, preferably from 0.7 to 6 g/10 minutes, elongation at break (ASTM D 638) from over 100% to 800%, preferably from 120 to 550%, and flexural modulus (ASTM D 790) from 100 to less than 3,000 MPa, preferably from 100 to 2700 MPa.

The solubility and insolubility of the polymers of the present invention are defined as fractions soluble or insoluble in xylene at ambient temperature, i.e., around 25°C (see note 1).

The above mentioned compositions can easily be applied to the exterior surface of metal pipes by using extrusion techniques known in the art, because they have a sufficiently high viscosity in the molten state, and therefore can withstand the tensions to which they are subjected during traditional coating processes without melt fracture.

In order to obtain an optimum adhesion to the pipe surface, it is preferable that prior to the application of the compositions of the invention said surface be coated with a layer of a hot melt composition of the type commonly used in the art (such as one based of polypropylene grafted with maleic anhydride).

Heterophasic composition (B) preferably has a propylene content in the copolymers of Fraction 1° ranging from 90 to 99% by weight. The polymer fraction insoluble in xylene in said

Fraction 1° more preferably ranges from 85 to 99% in the case of homopolymers, and from 85 to 95% in the case of copolymers.

Examples of the above mentioned  $C_4$ - $C_{10}$   $\alpha$ -olefins present in composition (B) are 1-butene, 1-pentene, 4-methyl-1-pentene,

1-hexene, and 1-octene. The preferred comonomer is 1-butene.

When present, the diene in composition (B) preferably ranges from 1 to 10%, more preferably 2.5 - 7% by weight with respect to the total weight of Fraction 2°. Examples of dienes are butadiene, 1,4-hexadiene, 1,5-hexadiene, and 5-ethylidene-2-norbornene.

When present, said Fraction (3°) preferably exceeds 1% by weight, more preferably ranging from 1 to 25% by weight with respect to the sum of Fractions (1°) and (2°). Preferably the content of ethylene in Fraction (3°) is at least 75% by weight, more preferably 80% by weight, with respect to the total weight of Fraction (3°). The comonomers in the copolymer of Fraction (3°) are preferably the same as those of the copolymer of Fraction (2°). An example of copolymer comprised in Fraction (3°) is an essentially linear semicrystalline copolymer of ethylene with propylene, a specific example of which is linear low density polyethylene (LLDPE).

The above mentioned heterophasic composition can be prepared by blending Fractions (1°), (2°), and optionally (3°) in the molten state, that is to say at temperatures greater than their softening or melting point, or more preferably by sequential polymerization in one or more stages in the presence of a highly stereospecific Ziegler-Natta catalyst. In particular, the catalyst system used comprises (a) a solid catalytic component containing a titanium compound and an electron-donor compound, both supported on magnesium chloride, and (b) an Altrialkyl compound and an electron-donor compound.

The above mentioned sequential polymerization process for the production of the heterophasic composition comprises at least two stages, where in the first stage the propylene is polymerized to form Fraction (1°), and in the subsequent stages mixtures of ethylene with propylene and/or said C<sub>4</sub>-C<sub>10</sub>α-olefin, and optionally diene, are polymerized to form Fraction (2°), and optionally Fraction (3°). The polymerization processes are carried out in liquid, gaseous, or liquid/gas phase. The reaction temperature in the various stages of polymerization can be equal or different, and generally ranges from 40° to 90°C, preferably from 50° to 80°C for the production of Fraction (1°), and from 40° to 60°C for the production of Fraction (2°) and optionally Fraction (3°).

Examples of sequential polymerization processes are described in European patent applications EP-A-472946 and EP-A-400333.

As a way of example, the heterophasic composition has MIL values ranging from 0.1 to 20 g/10 min, preferably from 0.2 to 15 g/10 min. The heterophasic composition with said melt index values can be obtained directly during the polymerization process; as an alternative, said heterophasic composition can be subjected to a chemical visbreaking process carried out in the presence of the appropriate visbreaking agents, such as peroxides. Said chemical visbreaking process is carried out according to well known methods.

Among the above mentioned compositions those that are particularly adequate for the preparation of the compositions of the present invention are the polyolefin compositions (B') comprising (weight percentage):

- 1) 30-50% of said Fraction (1°) (Fraction 1);
- 2) 50-70% of said Fraction (2°) where the quantity of ethylene in the copolymer is lower than 40%, preferably from 20 to 38% (Fraction 2); and

3) 0-20% of said Fraction (3°) (Fraction 3).

. Ev

In said composition (B') the weight percentage of the sum of fractions (2) and (3) with respect to the total composition ranges from 50 to 90%, preferably from 65 to 80%, and the weight ratio (3)/(2) is lower that 0.4, preferably ranging from 0.08 to 0.3.

As a way of example, the total quantity of ethylene in heterophasic composition (B') ranges from 15 to 35% by weight.

Other examples of the above mentioned compositions (B) that can be used in the present invention are the polyolefin compositions (B") comprising (weight percentage):

- I) 40-60% of said Fraction (1°) (Fraction I);
- II) 40-60% of said Fraction (2°) where the quantity of ethylene in the copolymer ranges from 40% to 70% (Fraction II); and
- III) 0-25% of said Fraction (3°) (Fraction III).

As a way of example, the total content of polymerized ethylene in said composition (B") ranges from 20 to 60% by weight.

The molecular weight of the various fractions of composition (B") (determined by measuring the intrinsic viscosity in tetrahydronaphtalene at 135°C) varies in function of the nature of the components, and the total melt index of the composition. In particular, the intrinsic viscosity is preferably comprised between the following limits:

- 0.5-3 dl/g for Fraction (I), and
- 2-8 dl/g for Fraction (II) plus Fraction (III).

Examples of heterophasic polyolefin compositions (B') and (B") are described respectively in the above mentioned European Patent applications EP-A-0 472946 and EP-A-400 333.

The above mentioned compatibilizers for fillers are preferably selected, for example, from stearic acid derivatives, such as inorganic stearates and stearamides, such as Ca and Zn stearates.

Further examples of fillers' compatibilizers different from the stearic acid derivatives are the polypropylene grafted with maleic anhydride or with other graftable compounds containing carboxylic groups, and systems based of vinyl-substituted alkoxysilanes and peroxides.

The compositions of the present invention can be prepared by using techniques known in the art for the preparation of polyolefin polymer blends with fillers. For example, one can use Banbury, Buss, or Brabender mixers at temperatures ranging from 180°C to 260°C.

The following examples are given in order to illustrate but not limit the present invention.

The methods used to obtain the data relative to the properties reported in the examples and the description are listed below.

Method Property ASTM-D 1238, condition L Melt Index (MIL) Solubility in xylene (see note 1 below) **ASTM D 412** Flexural Modulus (FM) **ASTM D 638** Tensile strength at 23°C **ASTM D 638** Yield stress at 23°C **ASTM D 638** Elongation at break at 23°C Elongation at yield at 23°C **ASTM D 638** B. S. 4370 Density Notched impact strength (Izod) **ASTM D 256** 

Note 1

7

Determination of percentage soluble in xylene: a solution is prepared of the sample in xylene at a concentration of 1% by weight, while the sample is maintained under agitation in xylene for one hour at 135°C. Continuing to stir, the content is allowed to cool to 95°C, after which the solution is poured into a 25°C bath, and left there for 20 minutes without stirring, and for 10 more minutes under stirring conditions. The solution is then filtered, and acetone is added to a portion of the filtrate in order to obtain the precipitation of the dissolved polymer. The polymer thus obtained is then recovered, washed, dried, and ultimately weighed in order to determine the percentage soluble in xylene.

#### Components used in the examples and comparative examples

- The following heterophasic compositions (B) (i)-(iii) are obtained by way of sequential polymerization in the presence of a high yield, highly stereospecific Ziegler-Natta catalyst supported on MgCl<sub>2</sub>.
  - Heterophasic composition (i) having a MIL of 0.6 g/10 min comprising (weight parts and percentages):
    - 1) 33 parts of a crystalline propylene random copolymer with 4.3% of ethylene; the copolymer containing about 9% of a fraction soluble in xylene at 25°C, and having an intrinsic viscosity [ $\eta$ ] of 1.5 dl/g;
    - 2) 61 parts of an ethylene/propylene amorphous copolymer containing 30% of ethylene, totally soluble in xylene at 25°C, and having an intrinsic viscosity [η] of 3.2 dl/g; and
    - 6 parts of an essentially linear ethylene/propylene copolymer totally insoluble in xylene at 25°C.

- Elastomeric composition (ii) having a MIL of 0.8 g/10 min comprising (weight parts and percentages):

- I. 40 parts of a crystalline propylene homopolymer having a xylene insoluble fraction of 97.5%, and MIL of 9 g/10 min;
- II. 47 parts of an amorphous ethylene/propylene copolymer, containing 50% of ethylene, totally soluble in xylene at 25°C, and having an intrinsic viscosity [n] of 3.7 dl/g; and
- III. 13 parts of an essentially linear ethylene/ propylene copolymer totally insoluble in xylene at 25°C.
- Elastomeric composition (iii) having a MIL of 10.5 g/10 min comprising (weight percentages):
  - 67% of a crystalline propylene random copolymer containing 3% of ethylene, having a MIL of 32 g/10 min;
  - 27% of an ethylene/propylene copolymer soluble in xylene at 25°C containing 55% of ethylene, and having an intrinsic viscosity [η] of 2.35 dl/g; and
  - 6% of an ethylene/propylene copolymer insoluble in xylene at 25°C.
- Polymer (iv): propylene random copolymer comprising (by weight) 3.5% of ethylene, and 6% of 1-butene, having a MIL of 5.5 g/10 min, and melt point of 132°C (determined by DSC).
- Polypropylenes (v) and (vi): crystalline propylene homopolymers containing 96% of a fraction insoluble in xylene at ambient temperature, and having a MIL of 6 and 10 g/10 min respectively.

Irganox B 225: 50% by weight of pentaerythritol-tetrakis[3(3,5-di-tert-butyl-4-hydroxyphenyl)]propionate (Irganox 1010) + 50% by weight of bis(2,4-di-tert-butylphenyl)phosphite (Irgafos 168), marketed by Ciba-Geigy.

#### Examples 1-5, and comparative examples 1c-3c

In a Buss 200 mixer are introduced the materials indicated in Table 1. The mineral filler is barium sulfate (sp. w = 4.4).

The above mentioned materials are extruded in the Buss at a temperature of 230°C.

Part of the composition thus obtained is used in an extrusion test using a single screw extruder with a flat die at a temperature of about 190°C, obtaining a homogeneous extrudate, and without melt fracture problems. The results prove that the composition is adequate for the use in pipe coating processes by way of extrusion.

In order to determine the mechanical properties, an other portion of the composition is used for the preparation of 120mm x 120mm x 3mm specimens by way of injection molding at 230°C.

The MIL value, density, and mechanical characteristics of the above mentioned compositions are reported in Table 2.

Table

Examples and comparative	_	2	۲	4	3	10	2c	3c
Barium sulfate %	76	75.5	80	85	75.5	75.5	75.5	9/
Elastom. Comp. (i) %	23.4	23.5	18.9	13.9	0	0	0	0
Elastom. Comp.(ii) %	0	0	0	0	23.5	0	. 0	0
Elastom. Comp.(iii)%	0	0	0	0	0	23.5	0	0
Polymer (iv) %	0	0	0	0	0	0	23.5	0
Polypropylene (v) %	0	0	0	0	0	0	0	21.5
Polypropylene (vi) %	0	0	0	0	0	0	0	2
Calcium Stearate %	0.5	0.5	8.0	0.8	0.5	0.5	0.5	0.5
Irganox B225 %	0.1	0.5	0.3	0.3	0.5	0.5	0.5	0

Table 2

Examples and comparative	ve 1	2	3	4	5	]C	2c	3c
examples								
MIL g/10 min	0.93	1.3	3.1	1.4	0.63	8.4	7.9	5.3
FM MPa	380	305	455	510	1020	1920	3710	2000
Yield stress %	4.7	4.3	4.8	4.9	6.4	9.2	12.8	•
Tensile strength %	,	7	,	1	•	-	•	-
Elongation at yield %	1	7	1	•	1	1	•	
Elongation at break %	440	450	375	355	220	10	20	45
Izod at 23°C J/m	NR <sup>13</sup>	NR¹)	NR¹)	NR")	452	252	<del>6</del> 4	•
Density g/ml	2.19	2.12	2.4	2.55	2.19	2.2	2.43	2.2

1) The specimen does not break.

#### **CLAIMS**

- 1. Polyolefin composition comprising:
  - (A) 60%-90% by weight of at least one heavy mineral filler with a specific weight greater than or equal to 3;
  - (B) 10-40% by weight of a heterophasic polyolefin composition; and
  - (C) a compatibilizer for fillers in quantities ranging from 0.1% to 4% by weight with respect to the weight of components (A) + (B);

said composition (B) being characterized in that it comprises (weight percentages):

- 1°) 30-60% of a propylene homopolymer fraction with an insolubility in xylene at ambient temperature greater than 80%, or a copolymer of propylene with ethylene and/or a C<sub>4</sub>-C<sub>10</sub> α-olefin, containing more than 80% of propylene, and having an insolubility in xylene at ambient temperature greater than 80% (Fraction 1°);
- 2°) 40-70% of a copolymer fraction of ethylene with propylene and/or a C<sub>4</sub>-C<sub>10</sub> α-olefin, and optionally minor quantities of diene; said copolymer containing from 15 to 70% of ethylene, and being soluble in xylene at ambient temperature (Fraction 2°); and
- 3°) 0-30%, with respect to the sum of Fractions (1°) and (2°), of a copolymer fraction containing ethylene, said fraction being insoluble in xylene at ambient temperature (Fraction 3°).
- Composition of claim 1 where the compatibilizer is selected from stearic acid derivatives.

3. Polyolefin composition of claim 1 where composition (B) comprises (weight percentages):

- 1) 30-50% of said Fraction (1°);
- 2) 50-70% of said Fraction (2°) where the quantity of ethylene in the copolymer is lower than 40%, preferably from 20 to 38%; and
- 3) 0-20% of said Fraction (3°).
- 4. Polyolefin composition of claim 1 where composition (B) comprises (weight percentages):
  - I) 40-60% of said Fraction (1°);
  - II) 40-60% of said Fraction (2°) where the quantity of ethylene in the copolymer ranges from 40% to 70%; and
  - III) 0-25% of said Fraction (3°).
- 5. Composition of claim 1 where the heavy mineral filler (A) is BaSO<sub>4</sub>, or Zr silicates.
- Use of the compositions of claim 1 for coating the outside surface of metal pipes by way of extrusion.
- 7. Metal pipes whose outside surface is coated with a layer of the compositions of claim 1.

## INTERNATIONAL SEARCH REPORT

inter nal Application No PCT/EP 98/06625

A. CLASSI IPC 6	FICATION OF SUBJECT MATTER C08K3/30 C08K3/34 C08L23	3/12 F16L58/16	
	Live and Debut Classification (IDC)	Western dupo	
	o International Patent Classification (IPC) or to both national clas SEARCHED	SITICATION AND IPC	
	ocumentation searched (classification system followed by classif	ication symbols)	
IPC 6	COSK COSL F16L		
Documental	tion searched other than minimum documentation to the extent the	hat such documents are included in the fields so	earched
Electronic d	ata base consulted during the international search (name of dat	a base and, where practical, search terms used	()
	·		
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of th	e relevant passages	Relevant to claim No.
X	DE 25 58 483 A (BASF AG) 7 July see page 4; claims; examples see page 5, paragraph 5	y 1977	1-5
А	EP 0 673 966 A (HIMONT INC) 27 September 1995 see page 2 - page 4		1-7
А	EP 0 254 375 A (WAVIN BV) 27 J see claims; examples	anuary 1988	1-5
		•	
Furt	ther documents are listed in the continuation of box C.	Patent family members are listed	d in annex.
	ategories of cited documents :	*T* later document published after the int	
consi	ent defining the general state of the art which is not dered to be of particular relevance document but published on or after the international	or priority date and not in conflict with cited to understand the principle or the invention "X" document of particular relevance; the	neory underlying the
which	date ent which may throw doubts on priority claim(s) or n is cited to establish the publication date of another on or other special reason (as specified)	cannot be considered novel or cannot involve an inventive step when the d "Y" document of particular relevance; the cannot be considered to involve an in-	ocument is taken alone claimed invention
"O" docum	nent referring to an oral disclosure, use, exhibition or means	document is combined with one or ments, such combination being obvin in the art.	nore other such docu-
later	ent published prior to the international filing date but than the priority date claimed	"&" document member of the same pater	<u></u>
Date of the	actual completion of the international search	Date of mailing of the international se	earch report
2	Prebruary 1999	11/02/1999	
Name and	mailing address of the ISA  European Patent Office, P.B. 5818 Patentlaan 2  NL - 2280 HV Rijswijk	Authorized officer	
	Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3018	Friederich, P	

## INTERNATIONAL SEARCH REPORT

Interi 1al Application No PCT/EP 98/06625

Patent do cited in sea			Publication date		tent family ember(s)	Publication date
DE 2558	3483	A	07-07-1977	NONE		<u> </u>
EP 0673	3966	A	27-09-1995	IT US AU AU BR CA JP NO	1269558 B 5494953 A 690200 B 1488795 A 9501158 A 2145107 A 8053586 A 951083 A	08-04-1997 27-02-1996 23-04-1998 28-09-1995 04-06-1996 23-09-1995 27-02-1996 25-09-1997
EP 0254	1375	A	27-01-1988	NL DE DK ES FI JP JP JP	8601910 A 3773568 A 8718020 U 385087 A 2027279 T 873213 A,B, 1902396 C 6031657 B 63092887 A 4957793 A	16-02-1988 14-11-1991 05-11-1992 24-01-1988 01-06-1992 24-01-1988 08-02-1995 27-04-1994 23-04-1988 18-09-1990